Charge when additional time is required to put into effect the corrective measures or to complete the investigation. The request for extension of time shall be accompanied by a progress report showing what has been accomplished to date.

§18.119 Importation.

ISM equipment shall be refused entry or withdrawal for consumption into the Customs territory of the United States, unless accompanied by a copy of FCC Form 740, in accordance with the provisions of subpart K, part 2 of this chapter.

§18.121 Exemptions.

Non-consumer ultrasonic equipment, and non-consumer magnetic resonance equipment, that is used for medical diagnostic and monitoring applications is subject only to the provisions of §§18.105, 18.109 through 18.119, 18.301 and 18.303 of this part.

[59 FR 39472, Aug. 3, 1994; 60 FR 47302, Sept. 12, 1995]

Subpart B—Applications and Authorizations

§18.201 Scope.

This subpart contains the procedures and requirements for authorization to market or operate ISM equipment under this part.

§18.203 Equipment authorization.

(a) Consumer ISM equipment, unless otherwise specified, shall be subject to certification prior to use or marketing. An application for certification shall be filed with the Commission on an FCC Form 731, pursuant to the relevant sections in part 2, subpart J of this chapter and shall also be accompanied by:

(1) A description of measurement facilities pursuant to §18.205 or reference to such information already on file with the Commission.

(2) A technical report pursuant to §§18.207 and 18.311.

NOTE: The Commission will accept applications for either type approval or certification until September 1, 1986, for consumer microwave ovens. After that date, only applications for certification will be accepted for filing.

(b) Consumer ultrasonic equipment generating less than 500 watts and operating below 90 kHz, and non-consumer ISM equipment shall be subject to verification, in accordance with the relevant sections of part 2, subpart J of this chapter.

(c) Grants of equipment authorization issued, as well as on-site certifications performed, before March 1, 1986, remain in effect and no further action is required.

§18.205 Description of measurement facilities.

(a) Any party filing a report of measurements with the Commission shall include in that report a description of the measurement facilities. If such a description is already on file with the Commission, it may be included by reference.

(b) The description shall include the following information:

(1) Location of test site.

(2) Physical description of the test site accompanied by photographs A4 (21 cm × 29.7 cm) or 8 × 10 inches (20.3 cm × 25.4 cm) in size. Photographs smaller than A4 (21 cm × 29.7 cm) or 8 × 10 inches (20.3 cm × 25.4 cm) will be acceptable if they are of sufficient clarity and mounted on A4 (21 cm × 29.7 cm) paper or paper 8 × 10 inches (20.3 cm × 25.4 cm).

(3) Scaled drawing showing the dimensions of the site, the physical layout of supporting structures and all structures within 5 times the distance between the measuring set and the device under test.

(4) Description of structures used to support the device being measured and the test instrumentation.

(5) List of measuring equipment used and information concerning the calibration of the measuring equipment, i.e., when the equipment was last calibrated and frequency of calibration.

(6) A statement indicating whether this facility is available to do measurement work for others on a contract basis.

(c) This information shall be kept current at all times. At least every three (3) years, the organization filing